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PTO/SB/21 (6-98)
Approved for use through 09/30/2000. OMB 0651-0031
Patent and Trademark Office: U.S. DEPARTMENT OF COMMERCE

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4		Application Number	09/945,152	
TRANSMITTAL	-	Filing Date	August 31, 2001	
FORM		First Named Inventor	Boyle et al.	
(to be used for all correspondence after initial	filing)	Group Art Unit		
		Examiner Name		
Total Number of Pages in This Submission	11	Attorney Docket Number	ACS-57082	

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ENCLOSURES (check all that apply)							
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Fee Attack			Drawing(s) g-related Papers		Щ	of Appeals and Interferences Appeal Communication to Group
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Extension of Tin	ne Request		Power of Change Address	Attorney, Revocation Correspondence	on	\mathbf{x}	Additional Enclosure(s) (please identify below):
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SIGNATURE OF APPLICANT, ATTORNEY, OR AGENT							
Firm <i>or</i> Individual name Ho	ward N. Som	mers;					
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PTO/SB/17 (10-01)
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U.S. Patent and Trademark Office; U.S. DEPARTMENT OF COMMERCE
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Patent fees are subject to annual revision.

TOTAL AMOUNT OF PAYMENT

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Complete if Known				
Application Number	09/945,152			
Filing Date	August 31, 2001			
First Named Inventor	Boyle et al.			
Examiner Name				
Group Art Unit				
Attorney Docket No.	ACC 57082			

METHOD OF PAYMENT	FEE CALCULATION (continued)					
The Commissioner is hereby authorized to charge indicated fees and credit any overpayments to:	3. ADDITIONAL FEES Large Small					
Deposit Account 06 27.25	Entity Entity					
Number 06-2425	Fee Fee Fee Fee Fee Description F	ee Paid				
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Charge Any Additional Fee Required Under 37 CFR 1.16 and 1.17	127 50 227 25 Surcharge - late provisional filing fee or cover sheet					
Applicant claims small entity status.	139 130 139 130 Non-English specification					
See 37 CFR 1,27	147 2,520 147 2,520 For filling a request for ex parte reexamination					
2. Payment Enclosed: Check Credit card Money Order Other	112 920* 112 920* Requesting publication of SIR prior to Examiner action					
FEE CALCULATION	113 1,840* 113 1,840* Requesting publication of SIR after Examiner action					
1. BASIC FILING FEE	115 110 215 55 Extension for reply within first month					
Large Entity Small Entity	116 400 216 200 Extension for reply within second month					
Fee Fee Fee Fee Description Code (\$) Code (\$) Fee Paid	117 920 217 460 Extension for reply within third month					
101 740 201 370 Utility filing fee	118 1,440 218 720 Extension for reply within fourth month					
106 330 206 165 Design filing fee	128 1,960 228 980 Extension for reply within fifth month					
107 510 207 255 Plant filing fee	119 320 219 160 Notice of Appeal					
108 740 208 370 Reissue filing fee	120 320 220 160 Filing a brief in support of an appeal					
114 160 214 80 Provisional filing fee	121 280 221 140 Request for oral hearing					
	138 1,510 138 1,510 Petition to institute a public use proceeding					
SUBTOTAL (1) (\$)	140 110 240 55 Petition to revive - unavoidable					
2. EXTRA CLAIM FEES	141 1,280 241 640 Petition to revive - unintentional					
Fee from Extra Claims <u>below</u> Fee Paid	142 1,280 242 640 Utility issue fee (or reissue)					
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104 280 204 140 Multiple dependent claim, if not paid	(37 CFR § 1.129(a))					
109 84 209 42 ** Reissue Independent claims over original patent	149 740 249 370 For each additional invention to be examined (37 CFR § 1.129(b))					
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**or number previously paid, if greater; For Reissues, see above	*Reduced by Besic Filing Fee Paid SUBTOTAL (3)					

SUBMITTED BY				Complete (#	applicable)
Name (Print/Type)	Howard N. Sommers	Registration No. (Attorney/Agent)	24,138	Telephone	310-824-5555
Signature	Howard. N. Sommers			Date	10/19/01

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PATENT Docket No. ACS-57082 (22272.3)

I hereby certify that this correspondence is being deposited with the United States Postal Service as First Class Mail in an envelope addressed to Commissioner for Patents, Washington, D.C. 20231, or. October 19, 2001.

Kroach N. Sommers

Howard N. Sommers, Registration No. 24,138 Date: October 19, 2001

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Application of) Examiner:
Inventor: Boyle et al.) Group Art Unit:
Serial Number: 09/945,152) Docket No. ACS-57082 (22272.3)
Filing Date: August 31, 2001) Date: October 19, 2001
For: SHEATHLESS EMBOLIC PROTECTION SYSTEM	
)

Commissioner for Patents Washington, D. C. 20231

PRE-EXAMINATION AMENDMENT

Dear Sir:

Please amend the above-identified application as follows:

IN THE SPECIFICATION

Please enter the following substitute paragraphs from the specification.

Please substitute page 8, line 24 - page 9, line 9, as follows:

In the drawings, wherein like reference numerals denote like or corresponding parts throughout the drawing figures, and particularly in the embodiments in accordance with the invention as shown in FIGS. 1-10, for example, a system 10 is provided for enabling an interventional procedure to be performed in a blood vessel 12 at an area of treatment 14. The system 10 is atraumatic, to inhibit injury to the patient. It includes a guide wire 16 which enables the system 10 to be positioned distal to the area of treatment 14. The system 10 is placed within the carotid artery 18 or other blood vessel of the patient, and is guided into position by the guide wire 16. The guide wire 16 includes a tip coil 20 at a distal end 22 thereof. The tip coil includes a proximal end 24. The tip coil 20 is attached at the proximal end thereof to the guide wire 16 for example by solder. The carotid artery 18 has the area of treatment 14 therein, which comprises the interventional procedure site, wherein atherosclerotic plaque 26 has built up against the inside wall 28, which decreases the diameter of the carotid artery 18. As a result, blood flow is diminished through this area.

Please substitute page 11, line 28 - page 12, line 10, as follows:

The system 10 further includes a delivery enabling element 82, which bears against the compressed filter device 30 for enabling delivery thereof to the position distal to the interventional procedure site 14, without extending about the filter device 30. The delivery enabling element 82 is also able to be withdrawn from bearing against the filter device 30. The delivery enabling element 82 includes an inner tube 84, which is extendable about the guide wire 16, and which includes a distal end 86 which is extendable into the filter device 30, through the channel 64 in the proximal portion 34 thereof, so as to bear against the compressing element 38. The inner tube 84 also pushes the tab members 74 radially outwardly and into engagement therewith

upon extending through the channel 64. The delivery enabling element 82 also includes an outer tube 88, extendable about the inner tube 84, which bears against the proximal portion 34 of the filter device 30 for delivery thereof.

Please substitute page 14, lines 8-17, as follows:

In the first version of the first embodiment of the present invention, as shown in FIGS. 1-5, the slots 80 in the engaging element 70 are engaged with the tab members 74 of the engageable element 68, to compress the filter device 30. An assembly of the compressed filter device 30 is inserted for example over the proximal end of the guide wire 16 extending outside the patient. The compressed filter device 30 is advanced over the proximal end of the guide wire 16 into the patient's body and onto the distal end 22 of the guide wire 16. The distal end 86 of the inner tube 84 of the delivery enabling element 82 is extended through the channel 64 in the proximal portion 34 of the filter device 30 so as to bear against the engaging element 70, to retain the filter device 30 in the compressed condition thereof. The outer tube 88 of the delivery enabling element 82 bears against the proximal portion 34 of the filter device 30 for enabling delivery of the filter device 30 to the location for deployment thereof. Delivery systems may be configured in over the wire or rapid exchange delivery platforms.

Please substitute page 14, lines 18-27, as follows:

Upon reaching the location distal to the interventional procedure site 14, the distal end 86 of the inner tube 84 is pulled in the proximal direction away from its position bearing against the engaging element 70, to a position for example extending slightly distal of the tabs 66, leaving a space between the distal end 86 of the inner tube 84 and the engaging element 70. The guide wire 16 is then pulled in the proximal

direction, pulling the stop member 72 into engagement with the engaging element 70. Upon pulling the guide wire 16 further in the proximal direction, the tab members 74 of the engageable element 68 slide out of the slot 80 in the engaging member 70, releasing the tab members 74 from the slots 80 so as to enable expansion and deployment of the filter device 30. Alternatively, for example, a slightly larger tip coil 20 may be used to push the engaging element 70 and deploy the filter device 30.

Please substitute page 14, line 28 - page 15, line 8, as follows:

The slots 90 of the inner tube 84, in the second version of the first embodiment of the invention, as depicted in FIGS. 6-8, engage the tab members 94 of the engageable element 68, to compress the filter device 30, and to retain the filter device 30 in the compressed condition during delivery. The outer tube 88 bears against the proximal portion 34 of the filter device 30 for enabling delivery of the filter device to the deployment location thereof. The distal end 86 of the inner tube 84 is pulled in the proximal direction, away from engagement with the engageable element 68, upon reaching the position distal to the interventional procedure site 14, for releasing the tab members 74 from the slots 80, and the tabs 66 engage the guide wire 16, for enabling expansion and deployment of the filter device 30.

Please substitute page 15, lines 9-27, as follows:

As illustrated in FIGS. 9-10, in the second embodiment of the present invention, an assembly of the filter device 30 and the obturator 40 is inserted for example over the proximal end of the guide wire 16 up to the position where the tabs 66 snap-fit into the space 100 so as to bear against the stop 98. The spring 92 is expanded, and the struts 102 of the filter device 30 engage the distal section 106 of the engageable element 68. The guide wire 16 is then pushed through the patient's